

GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

Run on: January 18, 2001, 04:20:59 ; Search time 1474.31 Seconds  
(without alignments)  
138.851 Million cell updates/sec

Title: PCT-US00-32259-12  
Perfect score: 40  
Sequence: 1 ctgagccaccataacccataactccaggattggg 40

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 1118133 seqs, 2558875100 residues  
Total number of hits satisfying chosen parameters: 2236266

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl:\*

- 1: gb\_bal:\*
- 2: gb\_ba2:\*
- 3: gb\_om:\*
- 4: gb\_ov:\*
- 5: gb\_ph:\*
- 6: gb\_pl1:\*
- 7: gb\_pl2:\*
- 8: gb\_pr1:\*
- 9: gb\_pr2:\*
- 10: gb\_pr3:\*
- 11: gb\_ro:\*
- 12: gb\_sy:\*
- 13: gb\_un:\*
- 14: em\_fun:\*
- 15: em\_hum1:\*
- 16: em\_hum2:\*
- 17: em\_in:\*
- 18: em\_om:\*
- 19: em\_or:\*
- 20: em\_ov:\*
- 21: em\_pat:\*
- 22: em\_ph:\*
- 23: em\_pl:\*
- 24: em\_ro:\*
- 25: em\_sts:\*
- 26: em\_sy:\*
- 27: em\_un:\*
- 28: em\_vi:\*
- 29: gb\_htg1:\*
- 30: gb\_htg2:\*
- 31: gb\_in1:\*
- 32: gb\_in2:\*
- 33: em\_ba1:\*
- 34: em\_ba2:\*
- 35: em\_hum3:\*
- 36: em\_hum4:\*
- 37: gb\_pr4:\*
- 38: gb\_htg3:\*
- 39: gb\_htg4:\*
- 40: gb\_htg5:\*
- 41: gb\_htg6:\*
- 42: gb\_htg7:\*
- 43: em\_htg1:\*

- 44: em\_htg2:\*
- 45: em\_htg3:\*
- 46: em\_hum5:\*
- 47: gb\_pl3:\*
- 48: gb\_pr5:\*
- 49: gb\_htg8:\*
- 50: gb\_htg9:\*
- 51: gb\_htg10:\*
- 52: gb\_htg11:\*
- 53: gb\_htg12:\*
- 54: gb\_htg13:\*
- 55: gb\_htg14:\*
- 56: gb\_in3:\*
- 57: gb\_htg15:\*
- 58: gb\_htg16:\*
- 59: gb\_htg17:\*
- 60: em\_htg4:\*
- 61: em\_htg5:\*
- 62: em\_htg6:\*
- 63: em\_htg7:\*
- 64: em\_hum6:\*
- 65: gb\_htg18:\*
- 66: gb\_htg19:\*
- 67: gb\_htg20:\*
- 68: gb\_htg21:\*
- 69: gb\_htg22:\*
- 70: gb\_htg23:\*
- 71: gb\_vil:\*
- 72: gb\_vil2:\*
- 73: gb\_ba3:\*
- 74: em\_htg8:\*
- 75: em\_htg9:\*
- 76: em\_htg10:\*
- 77: gb\_pr6:\*
- 78: gb\_pr7:\*
- 79: gb\_sts1:\*
- 80: gb\_sts2:\*
- 81: gb\_pat1:\*
- 82: gb\_pat2:\*
- 83: em\_htg0:\*
- 84: gb\_htg24:\*
- 85: gb\_pr8:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	% Match	Query Length	DB ID	Description
c	1	23	57.5	1653	37 AK026605	AK026605 Homo sapi
c	2	23	57.5	2600	78 HSTEGT	X75861 H.sapiens T
c	3	23	57.5	2634	81 AR058920	AR058920 Sequence
	4	23	57.5	2634	81 AR058931	AR058921 Sequence
	5	23	57.5	197189	42 AC019168	AC019168 Homo sapi
	6	22.8	57.0	171185	70 AP002424	AP002424 Homo sapi
	7	22.8	57.0	177097	70 AP001569	AP001569 Homo sapi
c	8	22.8	57.0	179726	9 AC007052	AC007052 Homo sapi
	9	22.8	57.0	188357	37 AL159996	AL159996 Human DNA
	10	22.8	57.0	200774	70 AP001592	AP001592 Homo sapi
	11	22.4	56.0	151514	77 HS471M13	Z97198 Human DNA s
	12	22	55.0	452	79 G22123	G22123 human STS W
c	13	22	55.0	34887	56 TBR012:99	AJ012199 Trypanoso
c	14	22	55.0	76727	77 HS821D:1	AL021453 Human DNA
c	15	22	55.0	110387	84 HSAJ96:1	AJ009611 Homo sapi
c	16	22	55.0	177540	9 AC006538	AC006538 Homo sapi
c	17	21.8	54.5	154599	41 AC016997	AC016997 Homo sapi
c	18	21.8	54.5	155907	49 AC020991	AC020991 Homo sapi
c	19	21.6	54.0	37470	48 CNS00YVD	AL096810 Homo sapi
c	20	21.6	54.0	104436	77 HS611N7	AL035663 Human DNA
c	21	21.6	54.0	168925	68 AL356502	AL356502 Homo sapi

```
c 22 21.6 54.0 208454 69 AL445199 Homo sapi
c 23 21.6 54.0 225653 69 AL359920 Homo sapi
c 24 21.2 53.0 70594 40 AC015786 Homo sapi
c 25 21.2 53.0 125219 69 AL445228 Homo sapi
c 26 21.2 53.0 147009 29 AC007618 Homo sapi
c 27 21.2 53.0 181215 49 AC021300 Homo sapi
c 28 21.2 53.0 208992 55 AC061709 Homo sapi
c 29 21.2 53.0 233454 9 AC005703 Homo sapi
c 30 21 52.5 2201 11 AF109155 Mus muscu
c 31 21 52.5 16332 1 AF104912 Escherich
c 32 21 52.5 80514 8 AC004454 Homo sapi
c 33 21 52.5 87235 50 AC022963 Homo sapi
c 34 21 52.5 92215 41 AC018277 Homo sapi
c 35 21 52.5 111269 30 AC009251 Drosophil
c 36 21 52.5 127409 37 AL137790 Human DNA
c 37 21 52.5 132571 69 AL445245 Homo sapi
c 38 21 52.5 155278 78 HSDJ655C4 Homo sapi
c 39 21 52.5 167277 52 AC025318 Homo sapi
c 40 21 52.5 167356 70 AP002814 Homo sapi
c 41 21 52.5 169480 65 AC079608 Homo sapi
c 42 21 52.5 173333 68 AC135590 Homo sapi
c 43 21 52.5 181233 38 AC105335 Homo sapi
c 44 21 52.5 186716 41 AC018468 Homo sapi
c 45 21 52.5 200627 42 AC019225 Homo sapi
```

ALIGNMENTS

```
RESULT 1
AK026605/c 1653 bp mRNA PRI 29-SEP-2000
LOCUS Homo sapiens cDNA: FLJ22952 fis, clone KAT09742.
AC026605
ACCESSION AK026605.1 GI:10439494
VERSION Homo sapiens signet-ring cell carcinoma cell_line:KATO III cDNA to
KEYWORDS oligo capping; fis (full insert sequence).
SOURCE mRNA, clone_lib:KAT clone:KAT09742.
ORGANISM Homo sapiens
```

```
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (sites)
Watanabe,K., Kumagai,A., Itakura,S., Yamazaki,M., Tashiro,H.,
Ota,T., Suzuki,Y., Obayashi,M., Nishi,T., Shibahara,T., Tanaka,T.,
Nakamura,Y., Isogai,T. and Sugano,S.
NEDO human cDNA sequencing project
Unpublished (2000)
2 (bases 1 to 1653)
Sugano,S., Suzuki,Y., Ota,T., Obayashi,M., Nishi,T., Isogai,T.,
Shibahara,T., Tanaka,T. and Nakamura,Y.
Direct Submission
Submitted (29-AUG-2000) to the DDBJ/EMBL/GenBank databases. Sumio
Sugano, Institute of Medical Science, University of Tokyo,
Laboratory of Genome Structure Analysis, Human Genome Center;
Shirokane-dai, 4-6-1, Minato-ku, Tokyo 108-8639, Japan
(E-mail:cdnal@ms.u-tokyo.ac.jp, Tel:81-3-5449-5286,
Fax:81-3-5449-5416)
```

```
COMMENT NEDO human cDNA sequencing project supported by Ministry of
International Trade and Industry of Japan; cDNA full insert
sequencing; Research Association for Biotechnology; cDNA library
construction, 5'- & 3'-end one pass sequencing; Department of
Virology and Human Genome Center, Institute of Medical Science,
University of Tokyo (partly supported by Science and Technology
Agency).
```

```
FEATURES Location/Qualifiers
source 1..1653
/organism="Homo sapiens"
/db_xref="taxon:9606"
/cell_line="KATO III"
/cell_type="signet-ring cell carcinoma"
/clone="KAT09742"
/clone_lib="KAT"
/note="cloning vector pME18SFL3"
```

```
BASE COUNT 441 a 369 c 358 g 485 t
ORIGIN
Query Match 57.5%; Score 23; DB 37; Length 1653;
Best Local Similarity 74.4%; Pred. No. 23;
Matches 29; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
Qy 2 tcgagccaccctcaaccctcaatactccagggattggg 40
||| ||||| ||| ||| ||| ||| ||| ||| ||| |||
Db 869 TCCACCCACACTTGACCCACAAAACATGCAGGGATTGGG 831
```

RESULT 2

```
HSTEGT/c HSTEGT 2600 bp mRNA PRI 28-SEP-1995
LOCUS HSTEGT H.sapiens TEGT gene.
DEFINITION H.sapiens TEGT gene.
ACCESSION X75861
VERSION X75861.1 GI:456258
KEYWORDS TEGT gene.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2600)
AUTHORS Walter,L.
TITLE Direct Submission
JOURNAL Submitted (14-JAN-1994) L. Walter, Abteilung Immunogenetik der
Universitaet Goettingen, Gosslerstr 12a, 37073 Goettingen, FRG
2 (bases 1 to 2600)
Walter,L., Marynen,P., Szpirer,J., Levan,G. and Gunther,E.
Identification of a novel conserved human gene, TEGT
Genomics 28 (2), 301-304 (1995)
MEDLINE 96015061
FEATURES Location/Qualifiers
source 1..2600
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="12q12-q13"
/clone_lib="lambda gt11 cDNA"
/tissue_type="testis"
41..754
/gene="TEGT"
41..754
/gene="TEGT"
/codon_start=1
/protein_id="CAA53472.1"
/db_xref="GI:458545"
/db_xref="SWISS-PROT:P55061"
/translation="MNIFDKINFDAALLAFSHITPSTQOHLKKVYASFALCMFVAAG
AVVWYTHFIQAGLLSALGSLIIMWIMATPHSHETEOKRLGGLAGFAFTLVGLGPGA
LEFCIAVNPISILPTAFWGTAMIFCTLSALYARRRSYFLGLGILMSALSLILSSIG
NVFFGSINWPFQANLYGLVYVMCGFVLVDQLIIIEKAHEGDODIYHWCIDFLDFITVF
RKULMILIANNEKDKKEKK"
```

```
BASE COUNT 619 a 621 c 590 g 770 t
ORIGIN
Query Match 57.5%; Score 23; DB 78; Length 2600;
Best Local Similarity 74.4%; Pred. No. 23;
Matches 29; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
Qy 2 tcgagccaccctcaaccctcaatactccagggattggg 40
||| ||||| ||| ||| ||| ||| ||| ||| ||| |||
Db 2024 TCCACCCACACTTGACCCACAAAACATGCAGGGATTGGG 1986
```

RESULT 3

```
AR058920/c AR058920 2634 bp DNA PAT 29-SEP-1999
LOCUS AR058920
DEFINITION Sequence 1 from patent US 58937838.
ACCESSION AR058920
VERSION AR058920.1 GI:5984497
```

## JOURNAL

Submitted (30-DEC-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
On Mar 13, 2000 this sequence version replaced gi:7021812.

## COMMENT

----- Genome Center -----  
Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: <http://genome.wustl.edu/gsc/index.shtml>  
----- Project Information -----  
Center project name: H\_NH0161E16  
----- Summary Statistics -----  
Sequencing vector: M13; 92%  
Chemistry: Dye-terminator; 3%  
Chemistry: Dye-terminator; 8% of reads  
Assembly: Program: Phrap; version 0.990319  
Consensus quality: 180505 bases at least Q40  
Consensus quality: 180445 bases at least Q30  
Insert size: 190000; agarose-fp  
Quality coverage: 3.69 in Q20 bases; agarose-fp  
Quality coverage: 3.65 in Q20 bases; sum-of-contigs

\* NOTE: This is a 'working draft' sequence. It currently consists of 24 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1451: contig of 1451 bp in length  
1551: gap of unknown length  
1552: contig of 1355 bp in length  
2907: gap of unknown length  
3006: contig of 1298 bp in length  
4304: gap of unknown length  
4404: contig of 1217 bp in length  
5621: gap of unknown length  
5721: contig of 2378 bp in length  
8095: gap of unknown length  
8199: contig of 2601 bp in length  
10800: gap of unknown length  
10900: contig of 3300 bp in length  
14201: gap of unknown length  
14301: contig of 3271 bp in length  
17571: gap of unknown length  
17671: contig of 4185 bp in length  
21856: gap of unknown length  
21957: contig of 3792 bp in length  
25749: gap of unknown length  
25849: contig of 3994 bp in length  
29843: gap of unknown length  
35421: contig of 5479 bp in length  
35522: gap of unknown length  
40974: contig of 5453 bp in length  
41075: gap of unknown length  
48591: contig of 7517 bp in length  
48691: gap of unknown length  
58168: contig of 9477 bp in length  
58268: gap of unknown length  
68032: contig of 9764 bp in length  
68132: gap of unknown length  
77463: contig of 9331 bp in length  
77563: gap of unknown length  
93029: contig of 15466 bp in length  
93129: gap of unknown length  
107954: contig of 14825 bp in length  
108054: gap of unknown length  
121842: contig of 13788 bp in length  
121942: gap of unknown length

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## FEATURES

## SOURCE

## BASE COUNT

## ORIGIN

## Query Match

## Best Local Similarity

## Matches

## Conservative

## Mismatches

## Indels

## Gaps

## 0;

## 0;

## 0;

## 0;

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```
* 121943 137362: contig of 15420 bp in length
* 137363 137462: gap of unknown length
* 137463 155524: contig of 18062 bp in length
* 155525 155624: gap of unknown length
* 155625 173416: contig of 17792 bp in length
* 173417 173517: gap of unknown length
* 173517 197189: contig of 23673 bp in length.
```

## FEATURES

```
source
  1..197189
    /organism="Homo sapiens"
    /db_xref="taxon:9606"
    /chromosome="12"
    /clone="RP11-161E16"
  1..1451
    /note="assembly_name:Contig3"
  1552..2906
    /note="assembly_name:Contig4"
  3007..4304
    /note="assembly_name:Contig5"
  4405..5621
    /note="assembly_name:Contig6"
  5722..8099
    /note="assembly_name:Contig7"
  8200..10800
    /note="assembly_name:Contig8"
  10901..14200
    /note="assembly_name:Contig9"
  14301..17571
    /note="assembly_name:Contig10"
  17672..21856
    /note="assembly_name:Contig11"
  21957..25748
    /note="assembly_name:Contig12"
  25849..29842
    /note="assembly_name:Contig13"
  29943..35421
    /note="assembly_name:Contig14"
  35522..40974
    /note="assembly_name:Contig15"
  41075..48591
    /note="assembly_name:Contig16"
  48692..58168
    /note="assembly_name:Contig17"
  58269..68032
    /note="assembly_name:Contig18"
  68133..77463
    /note="assembly_name:Contig19"
  77564..93029
    /note="assembly_name:Contig20"
  93130..107954
    /note="assembly_name:Contig21"
    clone_end:SP6
    vector_side:right
  108055..121842
    /note="assembly_name:Contig22"
  121943..137362
    /note="assembly_name:Contig23"
  137463..155524
    /note="assembly_name:Contig24"
  155625..173416
    /note="assembly_name:Contig25"
  173517..197189
    /note="assembly_name:Contig26"
  51312 a 47093 c 47437 g 49028 t 2319 others
ORIGIN
```

```
Query Match          57.5%  Score 23;  DB 42;  Length 197189;
Best Local Similarity 74.4%  Pred. No. 22;
Matches 29;  Conservative 0;  Mismatches 10;  Indels 0;  Gaps 0;
```

```
Oy  2 tggagccaccataacccctcaactccaggattggg 40
    || |||||  || ||||| || ||||| |||||
```

```
Db 123952 TCCACCCACACTTGAAGCCACAAAACCTGACGGGATTGGG 123990
```

## RESULT 6

```
AP002424
LOCUS      AP002424      171185 bp      DNA      HTG      31-MAY-2000
DEFINITION Homo sapiens chromosome 18 clone RP11-873L22 map 18q21, WORKING
            DRAFT SEQUENCE, in unordered pieces.
```

```
ACCESSION  AP002424.1 GI:8131688
```

```
VERSION    HTG; HTGS_PHASE1; HTGS_DRAFT.
```

```
KEYWORDS   Homo sapiens DNA, clone:RP11-873L22.
```

```
SOURCE     Homo sapiens
```

```
ORGANISM   Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
```

```
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
```

```
REFERENCE  1 (bases 1 to 171185)
```

```
AUTHORS    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
```

```
            Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
```

```
TITLE      Homo sapiens 171,185 genomic DNA of 18q21
```

```
JOURNAL    Published Only in DataBase (2000) In press
```

```
REFERENCE  2 (bases 1 to 171185)
```

```
AUTHORS    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
```

```
            Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
```

```
TITLE      Direct Submission
```

```
JOURNAL    Submitted (29-MAY-2000) to the DDBJ/EMBL/GenBank databases.
```

```
            Masahira Hattori, The Institute of Physical and Chemical Research
```

```
            (RIKEN), Genomic Sciences Center (GSC); Kitasato Univ., 1-15-1
```

```
            Kitasato, Sagamihara, Kanagawa 228-8555, Japan
```

```
            (E-mail:hattori@gsr.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
```

```
            Tel:81-42-778-9923, Fax:81-42-778-9924)
```

```
            ----- Genome Center
```

```
            Center: RIKEN Genomic Sciences Center (GSC)
```

```
            Center code: RIKEN
```

```
            Web site: http://hgp.gsc.riken.go.jp/
```

```
            Contact: hattori@gsr.riken.go.jp
```

```
            ----- Project Information
```

```
            Center project name: HumDrafit18
```

```
            Center clone name: RP11-873L22
```

```
            ----- Summary Statistics
```

```
            Sequencing vector: PCR products; 100% of reads
```

```
            Chemistry: Dye-terminator ET-amersham; 100% of reads
```

```
            Assembly program: Phrap; version 0.990329
```

```
            Consensus quality: 156977 bases at least Q40
```

```
            Consensus quality: 164110 bases at least Q30
```

```
            Consensus quality: 167421 bases at least Q20
```

```
            Insert size: 169485; sum-of-contigs
```

```
            Quality coverage: 4.80x in Q20 bases; sum-of-contigs
```

```
            -----
NOTE: This is a 'working draft' sequence. It currently consists of
18 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved
```

```
1
24923 contig of 24923 bp in length
47004 contig of 21981 bp in length
67070 contig of 19966 bp in length
82757 contig of 15587 bp in length
95544 contig of 12687 bp in length
109592 contig of 13948 bp in length
120167 contig of 10475 bp in length
129255 contig of 8988 bp in length
137003 contig of 7648 bp in length
144376 contig of 7273 bp in length
150659 contig of 6183 bp in length
155896 contig of 5137 bp in length
160522 contig of 4526 bp in length
163263 contig of 2641 bp in length
165194 contig of 1831 bp in length
167704 contig of 2410 bp in length
169750 contig of 1946 bp in length
171185 contig of 1335 bp in length
169851
```

\* NOTE: This is a 'working draft' sequence.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

```

FEATURES             Location/Qualifiers
     source            1..171185
                        /organism="Homo sapiens"
                        /db_xref="taxon:9606"
                        /chromosome="18"
                        /clone="RP11-873L22"
                        /map="18q21"
     misc_feature      1..24923
                        /note="assembly_fragment"
     misc_feature      25024..47004
                        /note="assembly_fragment"
     misc_feature      47105..67070
                        /note="assembly_fragment"
     misc_feature      67171..82757
                        /note="assembly_fragment"
     misc_feature      82858..95544
                        /note="assembly_fragment"
     misc_feature      95645..109592
                        /note="assembly_fragment"
     misc_feature      109693..120167
                        /note="assembly_fragment"
     misc_feature      120268..129255
                        /note="assembly_fragment"
     misc_feature      129356..137003
                        /note="assembly_fragment clone_end:SP6 vector_side:right"
     misc_feature      137104..144376
                        /note="assembly_fragment"
     misc_feature      144477..150659
                        /note="assembly_fragment"
     misc_feature      150760..155896
                        /note="assembly_fragment"
     misc_feature      155997..160522
                        /note="assembly_fragment"
     misc_feature      160623..163263
                        /note="assembly_fragment"
     misc_feature      163364..165194
                        /note="assembly_fragment clone_end:T7 vector_side:left"
     misc_feature      165295..167704
                        /note="assembly_fragment"
     misc_feature      167805..169750
                        /note="assembly_fragment"
     misc_feature      169851..171185
                        /note="assembly_fragment"
BASE COUNT  52031 a 32086 c 33289 g 52078 t 1701 others
ORIGIN
Query Match          57.0%; Score 22.8; DB 70; Length 171185;
Best Local Similarity 79.4%; Pred No. 27;
Matches 27; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Qy  1  ctgcagccaccataaccctcaactccaggg 34
    ||| ||| ||||| ||||| ||||| |||||
Db  93406  CTCTAGCTCCCGCCGCTTCATTCTCCAGTG 93439

RESULT  7
LOCUS      AP001569
DEFINITION Homo sapiens chromosome 18 clone RP11-859C31 map 18q21, WORKING
DRAFT SEQUENCE, in unordered pieces.
ACCESSION  AP001569
VERSION    AP001569.2 GI:8117403
KEYWORDS   HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE     Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 177097)

```

AUTHORS  
 TITLE  
 JOURNAL  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL

Hattori M., Ishii K., Toyoda A., Taylor T.D., Hong-Seog P.,  
 Fujiyama A., Yada T., Totoki Y., Watanabe H. and Sakaki Y.  
 Homo sapiens 177,097 genomic DNA of 18q21  
 Published Only in DataBase (2000) In press  
 2 (bases 1 to 177097)  
 Hattori M., Ishii K., Toyoda A., Taylor T.D., Hong-Seog P.,  
 Fujiyama A., Yada T., Totoki Y., Watanabe H. and Sakaki Y.  
 Direct Submission  
 Submitted (29-MAR-2000) to the DDBJ/EMBL/GenBank databases.  
 Masahira Hattori, The Institute of Physical and Chemical Research  
 (RIKEN), Genomic Sciences Center (GSC); Kitasato Univ., 1-15-1  
 Kitasato, Sagami-hara, Kanagawa 228-8555, Japan  
 (E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,  
 Tel:81-42-778-9923, Fax:81-42-778-9924)  
 On May 30, 2000 this sequence version replaced gi:7380904.  
 ----- Genome Center  
 Center: RIKEN Genomic Sciences Center (GSC)  
 Center code: RIKEN  
 Web site: http://hgp.gsc.riken.go.jp/  
 Contact: hattori@gsc.riken.go.jp  
 ----- Project Information  
 Center project name: HumTraff18  
 Center clone name: RP11-859C31  
 ----- Summary Statistics  
 Sequencing vector: PCR products; 100% of reads  
 Chemistry: Dye-terminator ET-amersham; 100% of reads  
 Assembly program: Phrap; version 0.990329  
 Consensus quality: 160670 bases at least Q40  
 Consensus quality: 168058 bases at least Q30  
 Consensus quality: 171631 bases at least Q20  
 Insert size: 173997; sum-of-contigs  
 Quality coverage: 4.63x in Q20 bases; sum-of-contigs

COMMENT

NOTE: This is a 'working draft' sequence. It currently consists of 32 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1	23792	contig cf	23792 bp in length
23893	45465	contig cf	21573 bp in length
45566	58167	contig cf	12602 bp in length
58268	71173	contig cf	12906 bp in length
71274	81253	contig cf	9980 bp in length
81354	91080	contig cf	9727 bp in length
91181	97387	contig cf	6207 bp in length
97488	104902	contig cf	7415 bp in length
105003	111107	contig cf	6105 bp in length
111208	116851	contig cf	5644 bp in length
116952	123585	contig cf	6634 bp in length
123686	128985	contig cf	5300 bp in length
129086	134233	contig cf	5148 bp in length
134334	138588	contig cf	4255 bp in length
138689	142573	contig cf	3885 bp in length
142674	146286	contig cf	3613 bp in length
146397	150053	contig cf	3667 bp in length
150154	154046	contig cf	3893 bp in length
154147	156225	contig cf	2079 bp in length
156326	158578	contig cf	2253 bp in length
158679	161124	contig cf	2446 bp in length
161225	163559	contig cf	2335 bp in length
163660	165538	contig cf	1879 bp in length
165639	166949	contig cf	1311 bp in length
167050	168240	contig cf	1191 bp in length
168341	169579	contig cf	1239 bp in length
169680	170978	contig cf	1299 bp in length
171079	172080	contig cf	1002 bp in length
172181	173514	contig cf	1334 bp in length
173615	174614	contig cf	1000 bp in length
174715	175976	contig cf	1262 bp in length
176077	177097	contig cf	1021 bp in length

Sequence updated (26-May-2000).

\* NOTE: This is a 'working draft' sequence.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

## FEATURES

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source
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    /db_xref="taxon:9606"
    /chromosome="18"
    /clone="RP11-859C21"
    /map="18q21"
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    /note="assembly_fragment"
  45566. .58167
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  91181. .97387
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  97488. .104902
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  111208. .116851
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  116952. .123585
    /note="assembly_fragment"
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  129086. .134233
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  134334. .138588
    /note="assembly_fragment"
  138689. .142573
    /note="assembly_fragment"
  142674. .146286
    /note="assembly_fragment clone_end:T7 vector_side:left"
  146387. .150053
    /note="assembly_fragment"
  150154. .154046
    /note="assembly_fragment"
  154147. .156225
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  168341. .169579
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  171079. .172080
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  173615. .174614
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  174715. .175976
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                  /note="assembly_fragment"
BASE COUNT      53531 a 33044 c 34236 g 53184 t 3102 others
ORIGIN
Query Match      57.0%; Score 22.8; DB 70; Length 177097;
Best Local Similarity 79.4%; Pred. No. 27;
Matches 27; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
Qy 1 ctcgagccatcccaataaccctcaatactccagg 34
    ||| |||| ||||| ||||| ||||| |||||
Db 3709 CTCAGGCTCCCATGCCCTCATTTTCAGTG 3742

RESULT 8
AC007052/c
LOCUS      AC007052 179726 bp DNA PRI 23-MAR-1999
DEFINITION Homo sapiens chromosome 18, clone hRPK.411_H_24, complete sequence.
ACCESSION  AC007052
VERSION    AC007052.4 GI:4510438
KEYWORDS   HTG.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 179726)
AUTHORS   Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE     Homo sapiens chromosome 18, clone hRPK.411_H_24
JOURNAL   Unpublished
REFERENCE  2 (bases 1 to 179726)
AUTHORS   Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
            Baker,J., Baldwin,J., Barna,N., Beckerly,R., Benn,J., Brown,A.,
            Castle,A., Cerny,J., Colangelo,M., Collins,S., Collymore,A.,
            Cooke,P., Dearellano,K., Depayre,E., Devon,K., Dewar,K.,
            Donelan,L., Doyle,M., Ferreira,P., FitzHugh,W., Forrest,C.,
            Funke,R., Gage,D., Galagan,J., Gardyna,S., Gilbert,D., Grant,G.,
            Hagos,B., Heaford,A., Horton,L., Howland,J.C., Jones,C., Kann,L.,
            Karatas,A., Lehoczy,J., Lieu,C., Locke,K., Macdonald,P.,
            Marquis,N., McEwan,P., McGurk,A., McKernan,K., McLaughlin,J.,
            Meldrim,J., Molla,M., Morris,W., Morrow,J., Mychaleckyj,J.,
            Naylor,J., Niloff,M., O'Connor,T., O'Donnell,P., Pavlin,B.,
            Peterson,K., Pollara,V., Riley,R., Roberts,D., Roy,A., Severy,P.,
            Stange-Thomann,N., Stojanovic,N., Stone,C., Subramanian,A.,
            Tesfaye,S., Torruella-Miller,I., Vassiliev,H., Vo,A., Wagner,A.,
            Wheeler,J., Wu,X., Wyman,D., Ye,W.J. and Zody,M.
            Direct Submission
TITLE     Submitted (09-MAR-1999) Whitehead Institute/MIT Center for Genome
JOURNAL   Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE  3 (bases 1 to 179726)
AUTHORS   Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
            Baker,J., Baldwin,J., Barna,N., Beckerly,R., Benn,J., Brown,A.,
            Castle,A., Cerny,J., Colangelo,M., Collins,S., Collymore,A.,
            Cooke,P., Dearellano,K., Depayre,E., Devon,K., Dewar,K.,
            Donelan,L., Doyle,M., Ferreira,P., FitzHugh,W., Forrest,C.,
            Funke,R., Gage,D., Galagan,J., Gardyna,S., Gilbert,D., Grant,G.,
            Hagos,B., Heaford,A., Horton,L., Howland,J.C., Jones,C., Kann,L.,
            Karatas,A., Lehoczy,J., Lieu,C., Locke,K., Macdonald,P.,
            Marquis,N., McEwan,P., McGurk,A., McKernan,K., McLaughlin,J.,
            Meldrim,J., Molla,M., Morris,W., Morrow,J., Mychaleckyj,J.,
            Naylor,J., Niloff,M., O'Connor,T., O'Donnell,P., Pavlin,B.,
            Peterson,K., Pollara,V., Riley,R., Roberts,D., Roy,A., Severy,P.,
            Stange-Thomann,N., Stojanovic,N., Stone,C., Subramanian,A.,
            Tesfaye,S., Torruella-Miller,I., Vassiliev,H., Vo,A., Wagner,A.,
            Wheeler,J., Wu,X., Wyman,D., Ye,W.J. and Zody,M.
            Direct Submission
TITLE     Submitted (23-MAR-1999) Whitehead Institute/MIT Center for Genome
JOURNAL   Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT   On Mar 24, 1999 this sequence version replaced gi:4432872.
            All repeats were identified using RepeatMasker: Smit, A.F.A. &
            Green, P. (1996-1997)
            http://ftp.genome.washington.edu/RM/RepeatMasker.html.
FEATURES   'Location/Qualifiers

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/note="AluY repeat: matches 3. .290 of consensus"
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/note="MLT1H repeat: matches 70. .532 of consensus"
31655. .32020
/note="THE1B repeat: matches 1. .364 of consensus"
32181. .32674
/note="match: GSS: Em:AQ822292"
32279. .32403
/note="MIR repeat: matches 57. .192 of consensus"
complement(32857. .33263)
33279. .33551
/note="match: GSS: Em:AQ139531"
35349. .35388
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35353. .35388
/note="10 copies 4 mer acac 92% conserved"
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/note="MIR repeat: matches 2. .246 of consensus"
36104. .36469
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/note="MLT2D repeat: matches 1. .553 of consensus"
38259. .38373
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36334. .38392
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38609. .38887
/note="match: STS: Em:G13414"
38781. .38991
/note="LTR33 repeat: matches 302. .518 of consensus"
39086. .39212
/note="LTR33 repeat: matches 29. .165 of consensus"
39249. .39483
/note="MER4D repeat: matches 1. .234 of consensus"
39483. .40142
/note="MER4D repeat: matches 359. .1017 of consensus"
40325. .40415
/note="MER94 repeat: matches 39. .134 of consensus"
41795. .41902
/note="LTR33 repeat: matches 370. .472 of consensus"
41903. .42192
/note="AluSg repeat: matches 1. .291 of consensus"
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/note="LTR33 repeat: matches 472. .521 of consensus"
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43695. .43862
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44083. .44384
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Query Match 57.0%; Score 22.8; DB 37; Length 188357;  
 Best Local Similarity 79.4%; Pred. NO. 27;  
 Matches 27; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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Oy 7 ccacccataaccctcaatactccaggattggg 40
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Db 56526 CCACCCCACTGTCTCTCAAGACTCCAGGGGAGGG 56559

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RESULT 10
AP001592
LOCUS
DEFINITION
Homo sapiens chromosome 18 clone RP11-869L2 map 18q21, WORKING
DRAFT SEQUENCE, in unordered pieces.
AP001592
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 200774)
AUTHORS
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE
Published Only in DataBase (2000) In press
JOURNAL
2 (bases 1 to 200774)
AUTHORS
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE
Direct Submission
JOURNAL
Submitted (31-MAR-2000) to the DDBJ/EMBL/GenBank databases.
Masahira Hattori, The Institute of Physical and Chemical Research
(RIKEN), Genomic Sciences Center (GSC); 1-15-1 Kitasato,
Sagamihara, Kanagawa 228-8555, Japan
(E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
Tel:81-42-778-9923, Fax:81-42-778-9924)
On Jul 31, 2000 this sequence version replaced gi:8117416.
COMMENT
----- Genome Center
Center: RIKEN Genomic Sciences Center (GSC)
Center code: RIKEN
Web site: http://hgp.gsc.riken.go.jp/
Contact: hattori@gsc.riken.go.jp
----- Project Information
Center project name: Humdraft18
Center clone name: RP11-869L2
----- Summary Statistics
Sequencing vector: PCR products; 100% of reads
Chemistry: Dye-terminator ET-amersham; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 197367 bases at least Q40
Consensus quality: 198735 bases at least Q30
Consensus quality: 199244 bases at least Q20
Insert size: 199774; sum-of-contigs
Quality coverage: 9.59x in Q20 bases; sum-of-contigs
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NOTE: This is a 'working draft' sequence. It currently consists of
11 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved.
1 36889 contig of 36889 bp in length
36990 73347 contig of 36358 bp in length
73448 98904 contig of 25457 bp in length
99005 124595 contig of 25591 bp in length
124696 145130 contig of 20435 bp in length
145231 163732 contig of 18502 bp in length
163833 181811 contig of 17979 bp in length
181912 190895 contig of 8984 bp in length
190996 198298 contig of 7303 bp in length
198399 199648 contig of 1250 bp in length
199749 200774 contig of 1026 bp in length.
* NOTE: This is a 'working draft' sequence.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
FEATURES
Location/Qualifiers
1. .200774
/organism="Homo sapiens"
/db_xref="taxon:9606"

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[illegible]

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TTEATKGIIDQAVQOKIDEAYIGERKIADEKAFGRKVFNAGGSQANCEGNIGE
NKASTLATLICQADNNGDTGSEHKACSGTAVTQWSGAANAPEOTVTYEMQLCD
TKDSHQITATQLFLEAVARQLRIINGAAYYGKRVAGNCNGEGGGLCVKTYDINN
AGKGFNSIPWDKLRQLREQLLEHERATKIEQNTALNCAAAATKALGRVQRREAA
GNSNAEPVVTQKSAKSEKQECNAAGDDPKKDLKGKCTYDEAKPKQKCTLSEED
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VSAFAALLF"
BASE COUNT 10433 a 6463 c 8050 g 9941 t
ORIGIN

Query Match 55.0%; Score 22; DB 56; Length 34887;
Best Local Similarity 73.7%; Pred. No. 60;
Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 3 cgaagcacccttaaccctcaatactccaggattggg 40
| ||||| ||| ||||| ||||| |||
Db 8362 CAAGCACCCAGACCATCAATCTCCACTGTATGG 8325

RESULT 14
HS821D11/c
LOCUS
DEFINITION
HS821D11 76727 bp DNA 12-DEC-1999
Human DNA sequence from clone RP5-821D11 on chromosome 22q12.3-13.1
Contains three partial unknown genes, one downstream of a predicted
CpG island, and the first coding exon of the SREBF2 gene for Sterol
Regulatory Element Binding transcription Factor 2 downstream of a
predicted CpG island, ESTs, STss, GSSs and genomic marker D22S1157,
complete sequence.
AL021453
ACCESSION
AL021453.1 GI:3413288
VERSION
HTG: D22S1157; SREBF2; Sterol Regulatory Element Binding
KEYWORDS
Transcription Factor 2.
human.
SOURCE
Homo sapiens
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
```

```
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
```

Mammalia: Eutheria; Primates; Catarrhini; Homidae; Homo.  
1 (bases 1 to 76727)  
Direct Submission  
Submitted (08-DEC-1999) Sanger Centre, Hinxton, Cambridgeshire,  
CB10 1SA, UK. E-mail enquiries: humqu@esanger.ac.uk  
requests: clonerequest@sanger.ac.uk  
On Aug 12, 1998 this sequence version replaced gi:3355590.  
This sequence has been finished according to sequence map criteria  
as follows. An attempt is made to resolve all sequencing problems,  
such as compressions and repeats, but not necessarily within known  
annotated human repeat sequence elements (e.g. Alu). Where the  
sequence is ambiguous, there is an annotation using the 'unsure'  
feature key.  
This sequence was generated from part of bacterial clone contigs of  
human chromosome 22, constructed by the Sanger Centre Chromosome 22  
Mapping Group. Further information can be found at  
http://www.sanger.ac.uk/HGP/Chr22  
During sequence assembly data is compared from overlapping clones.  
Where differences are found these are annotated as variations  
together with a note of the overlapping clone name. Note that the  
variation annotation may not be found in the sequence submission  
corresponding to the overlapping clone, as we submit sequences with  
only a small overlap as described above.  
The following abbreviations are used to associate primary accession  
numbers given in the feature table with their source databases:  
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information  
on the WORMPEP database can be found at  
http://www.sanger.ac.uk/Projects/C\_elegans/wormpep  
RP5-821D11 is from the library RPCI-5 constructed at the Roswell Park Cancer  
Institute by the group of Pieter de Jong. For further details see  
http://bacpac.med.buffalo.edu/  
VECTOR: pCYPAC2  
IMPORTANT: This sequence is not the entire insert of clone  
RP5-821D11 it may be shorter because we only sequence overlapping  
sections 'once, or longer because we arrange for a small overlap  
between neighbouring submissions.  
The true left end of clone RP5-821D11 is at 1 in this sequence. The  
true left end of clone CTA-250D10 is at 76628 in this sequence. The  
true right end of clone CTA-109G6 is at 42082 in this sequence. The  
start of this sequence overlaps with sequence Z99716 The end of  
this sequence overlaps with sequence Z83840.  
Location/Qualifiers  
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/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
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/clone="RP5-821D11"  
/clone\_lib="RPCI-5"  
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435..743  
/note="AluJb repeat: matches 8..312 of consensus"  
446..448  
/note="clone CTA-109G6  
tct in this entry  
substitution"  
/replace="ttt"  
744..840  
/note="L2 repeat: matches 2381..2490 of consensus"  
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/gene="dJ821D11.1"  
/note="match: ESTs: Em:AA419437"  
/evidence=not\_experimental  
/product="dJ821D11.1 (PUTATIVE protein)"  
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1314..1316  
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/note="clone CTA-109G6  
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substitution"

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CDS
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/note="other possible startcodon at 1583
this gene and dJ821D11.2 could be part of one gene
match: proteins: tr:O95505"
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/evidence=not_experimental
/product="dJ821D11.1 (PUTATIVE protein)"
/protein_id="CAAL6279.1"
/db_xref="GI:4200330"
/db_xref="SPTREMBL:O95505"
/translation="MLALTAKDSPRTALICSAWLLTASFSAQHKGSLQKDLLSQ
ACVGCLE"
variation
1760..1762
/gene="dJ821D11.1"
/note="clone CTA-109G6
gag in this entry
substitution"
1773..1923
/replace="ggg"
repeat_region
1901..1903
/note="MIR repeat: matches 47..201 of consensus"
variation
1901..1903
/gene="dJ821D11.1"
/note="clone CTA-109G6
tca in this entry
substitution"
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/replace="tta"
repeat_region
2010..2209
/note="12 repeat: matches 2250..2453 of consensus"
variation
2302..2304
/gene="dJ821D11.1"
/note="clone CTA-109G6
gga in this entry
substitution"
3042..3152
/replace="gaa"
repeat_region
3042..3152
/note="11M1 repeat: matches 6231..6332 of consensus"
3162..3457
/note="AluSp repeat: matches 1..297 of consensus"
3458..3551
/note="MIR repeat: matches 74..148 of consensus"
3552..3848
/note="AluDb repeat: matches 1..299 of consensus"
3849..3887
/note="MIR repeat: matches 32..74 of consensus"
4753..4754
/gene="dJ821D11.1"
/note="clone CTA-109G6
tt in this entry
deletion"
4772..4790
/replace="ttgagt"
repeat_region
4772..4790
/note="L1M4 repeat: matches 3167..3185 of consensus"
4791..5099
/note="AluSp repeat: matches 2..312 of consensus"
4791..4792
/gene="dJ821D11.1"
/note="clone CTA-109G6
tt in this entry
deletion"
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/replace="ttttatttat"
repeat_region
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5107..5112
/gene="dJ821D11.1"
/note="clone CTA-109G6
tttat in this entry
insertion"
5117..5427
/replace="tt"
repeat_region
5117..5427
/note="AluYb8 repeat: matches 1..318 of consensus"
5359..5361

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repeat_region
5430..5729
/note="AluSp repeat: matches 1..300 of consensus"
5730..5764
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5761..5762
/gene="dJ821D11.1"
/note="clone CTA-109G6
at in this entry
deletion"
5765..6075
/replace="att"
repeat_region
5765..6075
/note="AluY repeat: matches 1..311 of consensus"
5772..5773
/gene="dJ821D11.1"
/note="clone CTA-109G6
tt in this entry
deletion"
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variation
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/note="clone CTA-109G6
tt in this entry
deletion"
5780..5782
/replace="ttat"
variation
5780..5782
/gene="dJ821D11.1"
/note="clone CTA-109G6
tat in this entry
substitution"
5799..5801
/replace="ttt"
variation
5799..5801
/gene="dJ821D11.1"
/note="clone CTA-109G6
cgg in this entry
substitution"

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Query Match : 55.08; Score 22; DB 77; Length 76727;

Best Local Similarity 73.78; Pred. No. 59; Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ctccagccacccataaccctcaatactccaggatg 38

Db 34580 CTTGAGACACCAACCCCTCCGACACACAGGTTTG 34543

RESULT 15

LOCUS HSAJ9611/c

DEFINITION Homo sapiens chromosome 19 clone PAC RPCI-1 155110, \*\*\* SEQUENCING IN PROGRESS \*\*\*, 3 unordered pieces.

ACCESSION AJ009611

VERSION AJ009611.5 GI:9211527

KEYWORDS HTG: HTGS\_PHASE1; HTGS\_DRAFT.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 110387)

AUTHORS Radelof,U., Hennig,S., Ramser,J., Francis,F., Steffens,C.,

Klein,M., Seranski,P., Poustka,A., Reinhardt,R. and Lehrach,H.

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 110387)

AUTHORS MPIMG.

TITLE Direct Submission

JOURNAL Submitted (17-JUL-1998) MPIMG, Abt.Lehrach, Max Planck Institut

fuer Molekulare Genetik, Ihnestrasse 73, Berlin, 14195, Germany

COMMENT On Jul 15, 2000 this sequence version replaced gi:8248730.

contig 91 1..40399

```

contig 02 40500..61762
contig 03 61863..110387
Clone received from the Resource Centre of the Human Genome Project
at the Max-Planck-Institute for Molecular Genetics.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 40399: contig of 40399 bp in length
* 40400 40499: gap of 100 bp
* 40500 61762: contig of 21263 bp in length
* 61763 61862: gap of 100 bp
* 61863 110387: contig of 48525 bp in length.

FEATURES
    source
        1..110387
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /chromosome="19"
            /clone="PAC RPCI-1 155110"
            /clone_lib="RPCI1,3-5 Human PAC library, originating
            institute: Roswell Park Cancer Institute, creator: Pieter
            de Jong, P. Ioannou"
        misc_feature
            1 /note="T7_end:PAC RPCI-1 155110"
        misc_feature
            110387 /note="SP6_end:PAC RPCI-1 155110"
BASE COUNT 26958 a 27876 c 28291 g 27062 t 200 others
ORIGIN

```

```

Query Match 55.0%; Score 22; DB 84; Length 110387;
Best Local Similarity 73.7%; Fred. No. 59;
Matches 28; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 2 tcgagccaccataacctcaactccaggattgg 39
   ||| || |||| |||| |||| |||| |||| ||
DB 15347 TGGAGTCATCCCATGACCCCAAGACTCCAGTGCCAGG 15310

```

Search completed: January 18, 2001, 04:21:48  
Job time: 10474 sec

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GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 18, 2001, 11:04:24 ; Search time 262.85 Seconds  
(without alignments)  
57.168 Million cell updates/sec

Title: PCT-US00-32259-12

Perfect score: 40

Sequence: 1 ctcgagccaccataaacctcaactccaggattggg 40

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 480022 seqs, 187831343 residues

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N.Geneseq\_16.\*

- 1: /SID56/gcgdata/geneseq/geneseq/NA1980.DAT.\*
- 2: /SID56/gcgdata/geneseq/geneseq/NA1981.DAT.\*
- 3: /SID56/gcgdata/geneseq/geneseq/NA1982.DAT.\*
- 4: /SID56/gcgdata/geneseq/geneseq/NA1983.DAT.\*
- 5: /SID56/gcgdata/geneseq/geneseq/NA1984.DAT.\*
- 6: /SID56/gcgdata/geneseq/geneseq/NA1985.DAT.\*
- 7: /SID56/gcgdata/geneseq/geneseq/NA1986.DAT.\*
- 8: /SID56/gcgdata/geneseq/geneseq/NA1987.DAT.\*
- 9: /SID56/gcgdata/geneseq/geneseq/NA1988.DAT.\*
- 10: /SID56/gcgdata/geneseq/geneseq/NA1989.DAT.\*
- 11: /SID56/gcgdata/geneseq/geneseq/NA1990.DAT.\*
- 12: /SID56/gcgdata/geneseq/geneseq/NA1991.DAT.\*
- 13: /SID56/gcgdata/geneseq/geneseq/NA1992.DAT.\*
- 14: /SID56/gcgdata/geneseq/geneseq/NA1993.DAT.\*
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- 17: /SID56/gcgdata/geneseq/geneseq/NA1996.DAT.\*
- 18: /SID56/gcgdata/geneseq/geneseq/NA1997.DAT.\*
- 19: /SID56/gcgdata/geneseq/geneseq/NA1998.DAT.\*
- 20: /SID56/gcgdata/geneseq/geneseq/NA1999.DAT.\*
- 21: /SID56/gcgdata/geneseq/geneseq/NA2000.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	23	57.5	2634	19 V59067	Bax inhibitor BI-1
C 2	20.8	52.0	837	20 X37525	Human secreted pro
C 3	20	50.0	3364	19 V07266	Muscle creatine ki
C 4	19.8	49.5	2832	15 Q62186	Cyanobacteria repl
C 5	19.8	49.5	4809	15 Q62185	pAQL plasmid fragm
C 6	19.4	48.5	674	20 X37371	Human secreted pro
C 7	19.4	48.5	1554	20 X16675	Xenopus WA545 prot
C 8	19.2	48.0	300	20 Z14673	Human gene express
C 9	19.2	48.0	972	19 V37941	Rat arginase encod
C 10	19.2	48.0	1742	21 Z49765	Human DAP-kinase-r
C 11	19.2	48.0	1864	21 Z99730	CDNA encoding huma
C 12	19.2	48.0	2302	20 X86274	DNA encoding a hum

13	19	47.5	2401	14 Q50146	Restriction enzyme
14	18.8	47.0	1183	19 X14116	H. pylori GHPO 106
C 15	18.8	47.0	2191	21 Z92219	Human thiorodoxin
16	18.8	47.0	32207	20 V73805	KSHV LUR DNA (nucl
17	18.8	47.0	137507	19 V19941	KSHV long unique c
C 18	18.6	46.5	199	16 T13259	Human gene signatu
C 19	18.6	46.5	798	19 V30949	Human permidine/s
C 20	18.6	46.5	2532	20 Z52953	Human prostate tum
C 21	18.6	46.5	21170	20 X20535	Polynucleotide seq
C 22	18.4	46.0	453	20 V86886	EST clone B13. Ho
C 23	18.4	46.0	722	19 V28650	Ripening banana pu
C 24	18.4	46.0	1045	18 T59977	5' untranslated fe
C 25	18.4	46.0	1045	20 X02016	D. melanogaster ti
C 26	18.4	46.0	1194	21 Z91868	Streptococcus pneu
C 27	18.4	46.0	2082	15 Q55789	Sequence comprisin
C 28	18.4	46.0	2162	19 V37380	Streptococcus pneu
C 29	18.4	46.0	2382	12 Q10238	Encodes membrane e
C 30	18.4	46.0	2425	13 Q30133	ERR receptor gene.
C 31	18.4	46.0	2425	20 V59845	Murine ecotropic r
C 32	18.4	46.0	3954	18 T59975	Full length tipE p
C 33	18.4	46.0	3954	20 X02015	D. melanogaster ti
C 34	18.4	46.0	21338	19 V52153	Streptococcus pneu
C 35	18.2	45.5	259	17 T07319	Partial formyl-CoA
C 36	18.2	45.5	259	20 V64283	O. formigenes form
C 37	18.2	45.5	870	19 Z96201	S. pneumoniae deri
C 38	18.2	45.5	1233	20 Z08861	Human DNAX soluble
C 39	18.2	45.5	1281	20 X89654	CDNA encoding a hu
C 40	18.2	45.5	1305	20 Z06348	Nucleotide sequenc
C 41	18.2	45.5	1347	21 Z50347	Human orphan cytok
C 42	18.2	45.5	1391	19 V27144	Nucleotide sequenc
C 43	18.2	45.5	1498	21 Z34614	Human receptor mol
C 44	18.2	45.5	1579	19 V41689	Nucleotide sequenc
C 45	18.2	45.5	1579	20 X90753	Human U4 haematopo

ALIGNMENTS

RESULT 1	
ID V59067/C	V59067 standard; cDNA; 2634 BP.
XX	
AC V59067;	
XX	
DT 02-FEB-1999	(first entry)
XX	
DE Bax inhibitor BI-1 cDNA.	
XX	
KW Bax inhibitor; BI-1; human; apoptosis; ss.	
XX	
OS Homo sapiens.	
XX	
FH Key	Location/Qualifiers
FT CDS	73..786
FT	/*tag= a
XX	
PN WC9840397-A1.	
XX	
PD 17-SEP-1998.	
XX	
PF 13-MAR-1998;	98WO-US05015.
XX	
PR 14-MAR-1997;	97US-0818514.
XX	
PA (BURN-) BURNHAM INST.	
XX	
PI Reed JC, Xu Q;	
XX	
DR WPI: 1998-531519/45.	
DR P-PSDB; W73136.	
XX	
PT Bax inhibitor proteins, BI-1 and BI-2 - useful e.g. to modulate cellular apoptotic activity or identify agents altering BI-1 or BI-2	

PT binding which can modulate apoptotic activity  
 PS Claim 2; Page 61-63; 80pp; English.  
 XX  
 CC This cDNA clone codes for an inhibitor protein, termed BI-1 (see  
 CC W73136), of the pro-apoptotic protein Bax. Nucleic acids encoding  
 CC BI-1 and BI-2 (see V59068) were identified by suppression of  
 CC Bax-induced death of yeast cells transformed to express human Bax.  
 CC A human HepG2 cDNA library was used for library screening. The  
 CC invention provides vectors, optionally expression or viral vectors,  
 CC containing BI nucleic acids, and host cells containing these  
 CC vectors. The nucleic acids encoding BI-1/BI-2 can be used to  
 CC increase expression of these proteins in cells, or antisense  
 CC molecules prepared from them used to decrease expression. In  
 CC these ways, cellular apoptotic activity may be modulated (claimed).  
 CC The nucleic acids and complementary sequences are also useful as  
 CC probes to detect BI-encoding nucleic acid molecules in samples.  
 XX  
 SQ Sequence 2634 BP; 632 A; 626 C; 597 G; 779 T; 0 other;  
 Query Match 57.5%; Score 23; DB 19; Length 2634;  
 Best Local Similarity 74.4%; Pred. No. 0.92; Mismatches 0; Gaps 0;  
 Matches 29; Conservative 0; Indels 10; Indels 0; Gaps 0;  
 QY 2 tcgagccacccataaccctcaactccaggattggg 40  
 DB 2056 TCCACCCACTTGAAGCAGCACAAACTGCAGGGATTGG 2018  
 RESULT 2  
 X37525/c  
 ID X37525 standard; cDNA; 837 BP.  
 XX  
 AC X37525;  
 XX  
 DT 06-JUL-1999 (first entry)  
 XX  
 DE Human secreted protein cDNA fragment containing gene 75.  
 XX  
 KW Human; secreted protein; treatment; prevention; protein therapy; AIDS;  
 KW gene therapy; diagnosis; cancer; tumour; neurodegenerative disorder;  
 KW developmental abnormality; fetal deficiency; blood disorder; leukemia;  
 KW immune system disease; autoimmune disease; hepatic disease; lymphoma;  
 KW renal disease; inflammation; allergy; Alzheimer's disease; schizophrenia;  
 KW cognitive disorder; prostate disease; skeletal; cardiac; muscle disorder;  
 KW pulmonary disorder; transplant rejection; osteoclast; osteoporosis;  
 KW arthritis; malignancy; digestive; endocrine; infection; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN W09918208-A1.  
 XX  
 PD 15-APR-1999.  
 XX  
 PF 01-OCT-1998; 98WO-US20775.  
 XX  
 PR 02-OCT-1997; 97US-0060884.  
 PR 02-OCT-1997; 97US-0060833.  
 PR 02-OCT-1997; 97US-0060836.  
 PR 02-OCT-1997; 97US-0060837.  
 PR 02-OCT-1997; 97US-0060838.  
 PR 02-OCT-1997; 97US-0060839.  
 PR 02-OCT-1997; 97US-0060843.  
 PR 02-OCT-1997; 97US-0060862.  
 PR 02-OCT-1997; 97US-0060866.  
 PR 02-OCT-1997; 97US-0060874.  
 XX  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 PI Carter KC, Duan DR, Endress GA, Feng P, Ferrie AM;  
 PI Florence KA, Greene JM, Janat F, Lafleur DW, Ni J;  
 PI Rosen CA, Ruben SM, Shi Y, Young P, Yu G;

XX WPI; 1999-264022/22.  
 DR P-PSDB; Y07926.  
 XX  
 PT New isolated human genes and the secreted polypeptides they encode  
 XX  
 PS Claim 1a; Page 264; 368pp; English.  
 CC  
 CC This invention describes novel isolated human genes and the secreted  
 CC proteins they encode. The products of the invention are useful for  
 CC preventing, treating or ameliorating medical conditions, e.g. by protein  
 CC or gene therapy. Also pathological conditions can be diagnosed by  
 CC determining the amount of the new polypeptides in a sample or by  
 CC determining the presence of mutations in the new polynucleotides.  
 CC Specific uses are described for each of the 101 polynucleotides, based on  
 CC which tissues they are most highly expressed in, and include developing  
 CC products for the diagnosis or treatment of cancer, tumours,  
 CC neurodegenerative disorders, developmental abnormalities and fetal  
 CC deficiencies, blood disorders, leukemias, diseases of the immune system,  
 CC autoimmune diseases, hepatic and renal diseases, lymphomas, inflammation,  
 CC allergies, Alzheimer's and cognitive disorders, schizophrenia, prostate  
 CC disease, skeletal or cardiac muscle disorders, pulmonary disorders,  
 CC transplant rejection, disorders involving osteoclasts such as  
 CC osteoporosis, arthritis or malignancies, digestive/endocrine disorders,  
 CC infections and AIDS. The human secreted proteins of the invention are  
 CC represented in Y07852-Y07993 and the encoding nucleic acids are  
 CC represented in X37451-X37552.  
 XX  
 SQ Sequence 837 BP; 171 A; 249 C; 187 G; 228 T; 2 other;  
 Query Match 52.0%; Score 20.8; DB 20; Length 837;  
 Best Local Similarity 70.0%; Pred. No. 6.5;  
 Matches 28; Conservative 0; Mismatches 12; Indels 0; Gaps 0;  
 QY 1 ctgagccacccataaccctcaactccaggattggg 40  
 DB 250 CTCAGGCAGGCCAGAACACTCCAGAGCGCTGGAGATTGG 211  
 RESULT 3  
 V07266/c  
 ID V07266 standard; DNA; 3364 BP.  
 XX  
 AC V07266;  
 XX  
 DT 28-SEP-1998 (first entry)  
 XX  
 DE Muscle creatine kinase enhancer/promoter sequence.  
 XX  
 KW Encapsidated adenovirus minichromosome; EAM; gene therapy;  
 KW vector; muscle creatine kinase; promoter; mouse; dystrophin;  
 KW muscular dystrophy; ds.  
 XX  
 OS Mus sp.  
 XX  
 PN Key Location/Qualifiers  
 FT enhancer 2102..2307  
 FT prim\_transcript 3358..3364  
 FT /tag= a  
 FT /tag= b  
 FT /note= "5' end of primary transcript"  
 XX  
 PN W09817783-A1.  
 XX  
 PD 30-APR-1998.  
 XX  
 PF 23-OCT-1997; 97WO-US19541.  
 XX  
 PR 23-OCT-1996; 96US-0735609.  
 XX  
 PA (UNMI ) UNIV MICHIGAN.

PI Amalfitano A, Chamberlain JS, Hartigan-O'Connor DJ;  
 PI Hauser MA, Kumar-Singhr;  
 XX  
 DR WPI; 1998-261485/23.

XX New adenoviral recombinant plasmid(s) - comprise sequences provided  
 PT for expression of large foreign DNA fragments, used for, e.g. gene  
 PT therapy of genetic disease(s)

XX Example 6a; Page 105-107; 139pp; English.

XX This nucleotide sequence comprises the promoter/enhancer region  
 CC of the murine muscle creatine kinase (MCK) gene. The invention  
 CC provides encapsidated adenovirus minichromosomes (EAMs) containing  
 CC a full-length murine dystrophin cDNA. The EAM consists of an  
 CC infectious encapsidated linear genome containing adenovirus  
 CC origins of replication, packaging signal elements, a  
 CC beta-galactosidase reporter gene cassette and a full-length  
 CC (14 kb) dystrophin cDNA regulated by the MCK enhancer/promoter.  
 CC EAMs are generated by cotransfecting 293 cells with supercoiled  
 CC plasmid DNA (pAd5 beta-dys) containing an embedded inverted  
 CC origin of replication (and the remaining above elements) with  
 CC linear DNA from E1-deleted virions expressing human placental  
 CC alkaline phosphatase (hpAP). All proteins necessary for the  
 CC generation of EAMs are provided in trans from the hpAP virions.  
 CC The EAMs are useful for gene transfer to a variety of cell  
 CC types in vitro and in vivo. The invention provides improved  
 CC adenovirus vectors (see also V07261) and packaging cell lines  
 CC useful for a wide variety of gene therapy applications.

XX Sequence 3364 BP; 860 A; 900 C; 810 G; 794 T; 0 other;

Query Match 50.0%; Score 20; DB 19; Length 3364;  
 Best Local Similarity 72.2%; Pred. No. 18;  
 Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

OY 5 agcaccaccataaacctcaatactccaggattggg 40  
 ||||| ||| ||||| || ||||| |||  
 Db 2326 AGCCTGCCCTCACCCTGGATCCACGAGGACAGG 2291

RESULT 4  
 Q62186  
 ID Q62186 standard; DNA; 2832 BP.

XX AC Q62186;

XX 22-NOV-1994 (first entry)

XX Cyanobacteria replication protein coding sequence.

XX Plasmid; pAQ1; Cyanobacteria; PCC7002; vector; expression;  
 KW carbon dioxide; air; ss.

XX Agmenellum quadruplicatum PR-6.

XX JP0607877-A.

XX 22-MAR-1994.

XX 02-SEP-1992; 92JP-0274768.

XX 02-SEP-1992; 92JP-0274768.

XX (KANT ) KANSAI DENRYOKU KK.

XX WPI; 1994-131278/16.

XX P-PSDB; R51499.

XX New vector from cyanobacteria. - used for foreign gene expression  
 PT in cyanobacteria

XX

PS Disclosure; Page 15-18; 23pp; Japanese.

XX This sequence represents a fragment of the plasmid pAQ1 and encodes  
 CC a protein which has replication activity in Cyanobacteria. This  
 CC sequence was used in the production of a vector for the expression  
 CC of heterologous proteins in Cyanobacteria. This allows effective use  
 CC of carbon dioxide in the air.

XX Sequence 2832 BP; 760 A; 715 C; 724 G; 633 T; 0 other;

Query Match 49.5%; Score 19.8; DB 15; Length 2832;  
 Best Local Similarity 69.2%; Pred. No. 21;  
 Matches 27; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

OY 2 tcgagccaccataaacctcaatactccaggattggg 40  
 ||||| ||| || ||||| || ||||| |||  
 Db 305 tcgagctaccagaacgcgatcaaggcttcgggaatggg 343

RESULT 5  
 Q62185/C  
 ID Q62185 standard; DNA; 4809 BP.

XX AC Q62185;

XX 22-NOV-1994 (first entry)

XX pAQ1 plasmid fragment.

XX Plasmid; pAQ1; Cyanobacteria; PCC7002; vector; expression;  
 KW carbon dioxide; air; ss.

XX Agmenellum quadruplicatum PR-6.

XX JP06078777-A.

XX 22-MAR-1994.

XX 02-SEP-1992; 92JP-0274768.

XX 02-SEP-1992; 92JP-0274768.

XX (KANT ) KANSAI DENRYOKU KK.

XX WPI; 1994-131278/16.

XX New vector from cyanobacteria. - used for foreign gene expression  
 PT in cyanobacteria

XX Disclosure; Page 9-14; 23pp; Japanese.

XX This sequence represents a fragment of the plasmid pAQ1. This  
 CC plasmid was prepared from Cyanobacteria PCC7002. This sequence was  
 CC used in the production of a vector for the expression of heterologous  
 CC proteins in Cyanobacteria. This allows effective use of carbon  
 CC dioxide in the air.

XX Sequence 4809 BP; 1136 A; 1193 C; 1165 G; 1315 T; 0 other;

Query Match 49.5%; Score 19.8; DB 15; Length 4809;  
 Best Local Similarity 69.2%; Pred. No. 24;  
 Matches 27; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

OY 2 tcgagccaccataaacctcaatactccaggattggg 40  
 ||||| ||| || ||||| || ||||| |||  
 Db 2348 TCGAGCTACCAAGAACCGATCAGGCTTCTGGGAATGGG 2310

RESULT 6  
 X37371/c  
 ID X37371 standard; cDNA; 674 BP.

XX AC x37371;  
XX DT 06-JUL-1999 (first entry)  
XX DE Human secreted protein cDNA fragment containing gene 3.  
XX KW Human; secreted protein; prevention; treatment; protein therapy;  
XX KW gene therapy; diagnosis; cancer; tumour; neurodegenerative disorder;  
XX KW developmental abnormality; foetal deficiency; blood disorder; lymphoma;  
XX KW leukemia; immune system disorder; autoimmune disease; hepatic disease;  
XX KW renal disease; inflammation; allergy; asthma; sepsis; diabetes; AIDS;  
XX KW Alzheimer's disease; cognitive disorder; schizophrenia; osteoporosis;  
XX KW arthritis; psoriasis; digestive; endocrine; infection; ss.  
XX OS Homo sapiens.  
XX PN WO9909155-A1.  
XX PD 25-FEB-1999.  
XX PF 18-AUG-1998; 98WO-US17044.  
XX PR 16-JUN-1998; 98US-0092956.  
XX PR 15-JUL-1998; 98US-0092956.  
XX PR 19-AUG-1997; 97US-0056368.  
XX PR 19-AUG-1997; 97US-0056369.  
XX PR 19-AUG-1997; 97US-0056535.  
XX PR 19-AUG-1997; 97US-0056535.  
XX PR 19-AUG-1997; 97US-0056535.  
XX PR 19-AUG-1997; 97US-0056536.  
XX PR 19-AUG-1997; 97US-0056628.  
XX PR 19-AUG-1997; 97US-0056629.  
XX PR 19-AUG-1997; 97US-0056726.  
XX PR 19-AUG-1997; 97US-0056728.  
XX PA (HUMA-) HUMAN GENOME SCI INC.  
XX PI Brewer LA, Duan R, Ebner R, Endress GA, Feng P;  
XX PI Florence C, Florence KA, Komatsoulis GA, Lafleur DW;  
XX PI Moore PA, Olsen HS, Rosen CA, Ruben SM, Shi Y, Soppet DR;  
XX PI Young PE;  
XX WPI; 1999-190160/16.  
XX P-PSDB; Y07746.  
XX PT New isolated human genes and the secreted polypeptides they encode  
XX PT - useful for diagnosis and treatment of e.g. cancers, neurological  
XX PT disorders, immune diseases, inflammation or blood disorders  
XX PS Claim 1a; Page 192; 280pp; English.  
XX CC This invention describes novel isolated human secreted proteins and  
XX CC their encoding nucleic acid sequences. The products of the invention  
XX CC are useful for preventing, treating or ameliorating medical conditions  
XX CC e.g. by protein or gene therapy. Also pathological conditions can be  
XX CC diagnosed by determining the presence or amount of expression of  
XX CC the new polypeptides in a sample or by determining the presence or  
XX CC absence of mutations in the new polynucleotides. Specific uses are  
XX CC described for each of the 70 polynucleotides, based on which tissues they  
XX CC are most highly expressed in, and include developing products for the  
XX CC diagnosis or treatment of cancer, tumours, neurodegenerative  
XX CC disorders, developmental abnormalities and foetal deficiencies, blood  
XX CC disorders, leukemias, diseases of the immune system, autoimmune diseases,  
XX CC hepatic and renal disease, lymphomas, inflammation, allergies, asthma,  
XX CC sepsis, diabetes, Alzheimer's and cognitive disorders, schizophrenia,  
XX CC osteoporosis, arthritis, psoriasis, digestive/endocrine disorders,  
XX CC infections and AIDS. The human secreted proteins of the invention are  
XX CC represented in Y07744-Y07850 and the encoding nucleic acids are  
XX CC represented in X37369-X37441.  
XX SQ Sequence 674 BP; 238 A; 112 C; 175 G; 145 T; 4 other;

Query Match ; 48.5%; Score 19.4; DB 20; Length 674;  
Best Local Similarity 70.3%; Pred. No. 24;  
Matches 26; Conservative 0; Mismatches 11; Indels 0; Gaps 0;  
QY 4 gagccaccataaccctcaataactccaggattggg 40  
DB 645 GAGTCCTCCCACTCACCTTCAATACAAATGCATGGG 609  
RESULT 7  
X16675/c  
ID X16675 standard; DNA; 1554 BP.  
XX AC X16675;  
XX DT 29-APR-1999 (first entry)  
XX DE Xenopus WAS45 protein encoding DNA.  
XX KW Xenopus; WAS45 protein; TGF-beta; transforming growth factor beta;  
XX KW proliferation; differentiation; mesodermal tissue; neural; muscle;  
XX KW bone; cartilage; connective tissue; wound healing; gene therapy; ss.  
XX OS Xenopus sp.  
XX FH Key Location/Qualifiers  
XX CDS 55..1119  
FT /tag= a  
FT /product= "WAS45 protein"  
FT sig\_peptide 55..774  
FT /tag= b  
FT mat\_peptide 775..1116  
FT /tag= c  
XX PN WO9902678-A1.  
XX PD 21-JAN-1999.  
XX PF 24-APR-1998; 98WO-US08334.  
XX PR 10-JUL-1997; 97US-0890918.  
XX (GEMY ) GENETICS INST INC.  
XX (WHEED ) WHITEHEAD INST BIOMEDICAL RES.  
XX PI Lavallie ER, Kacie LA, Sive H, Sun B;  
XX WPI; 1999-120879/10.  
XX P-PSDB; W94656.  
XX PT New isolated Xenopus WAS45 DNA - which is a member of the  
XX PT transforming growth factor-beta superfamily, used to induce tissue  
XX PT formation and in wound healing  
XX PS Claim 1; Page 51-52; 73pp; English.  
XX CC The present sequence encodes the Xenopus WAS45 protein. WAS45 proteins  
XX CC have the ability to induce, enhance and/or inhibit the formation,  
XX CC growth, proliferation, differentiation, maintenance of mesodermal  
XX CC tissue, including neurons and/or related neural cells and tissues such  
XX CC as brain cells, Schwann cells, glial cells and astrocytes, as well as  
XX CC muscle cells and tissues. They can be used for treating bone, cartilage,  
XX CC muscle, nerve, epidermis and/or other connective tissue defects, as well  
XX CC as periodontal disease and healing of various epidermis, nerve, smooth  
XX CC including spinal cord, muscle, including cardiac, striated or smooth  
XX CC muscle, and other tissues and wounds, and other organs such as liver,  
XX CC pancreas, spleen, brain, lung, cardiac and kidney tissue. They can also  
XX CC be used to treat or prevent such conditions as osteoarthritis,  
XX CC osteoporosis, and other abnormalities of bone, cartilage, muscle, nerve,  
XX CC epidermis or other connective tissue, organs such as liver, pancreas,  
XX CC spleen, lung, cardiac and kidney and other tissues. They can also be  
XX CC used for wound healing, reduction of fibrosis and reduction of scar  
XX CC tissue formation. They can also be used to induce bone and/or cartilage

The present invention describes a library of human polynucleotides comprising the sequences given in Z12532 to Z17779. Also described is a method of detecting differentially expressed genes correlated with the cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample from a cell suspected of being cancerous, where the gene product is encoded by one of the 5248 polynucleotide sequences given in Z12532 to Z17779. The polynucleotides can be used as a source of primers and probes, which can be used for a variety of purpose, e.g. detection of expression levels,

Claim 13: page 54-66: 92pp: Japanese.

The present sequence encodes rat arginase. The present invention describes a tissue fibrosis inhibitor composition which contains as an active component an enzyme which can decompose arginine, the enzyme comprises human or rat arginase (e.g. derived from liver tissue) or arginine dehydase or their modified derivatives (such as polyethylene glycol-modified enzyme). The invention can be used in the prevention and treatment of fibrosis in tissues such as lung, kidney, liver, skin



```
Matches 27; Conservative 0; Mismatches 13; Indels 0; Gaps 0;
Qy 1 ctcgagccaccataaccctcaatactccaggattggg 40
   ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 955 ctcgagcattcccaagctcttaattctccataaaatggg 994

RESULT 12
X86274/c
ID X86274 standard; DNA; 2302 BP.
XX
AC X86274;
XX
DT 22-SEP-1999 (first entry)
XX
DE DNA encoding a human PIG protein.
XX
KW p53 transcription tag; p53 status; cancer; cytotoxicity; carcinogenicity;
KW neoplastic; PIG; ss.
XX
OS Homo sapiens.
XX
PN W09914356-A2.
XX
PD 25-MAR-1999.
XX
PF 17-SEP-1998; 98WO-US19300.
XX
PR 30-MAR-1998; 98US-0079817.
PR 17-SEP-1997; 97US-0059153.
XX
PA (UYJO ) UNIV JOHNS HOPKINS.
XX
PI Kinzler KW, Polyak K, Vogelstein B;
XX
DR WPI; 1999-443793/37.
XX
PT Use of p53 transcription tags to determine p53 status in, e.g.
PT cancer diagnosis
XX
PS Disclosure; Page 65-66; 73pp; English.
XX
CC The specification describes the use of p53 transcription tags for
CC developing products to determine p53 status, to diagnose cancer
CC and to evaluate cytotoxicity or carcinogenicity of a test agent.
CC A method for diagnosing cancer or determining p53 status in a sample
CC suspected for being neoplastic comprises comparing the level of
CC transcription of an RNA transcript in a first sample (s1) of a first
CC tissue (t1) to the level of transcription of the transcript in a second
CC sample (s2) of a second tissue (s2), where s1 is suspected of being
CC neoplastic and s2 is a normal human tissue (of the same type) and the
CC transcript is identified by a tag; and categorizing s1 as neoplastic
CC or as having a mutant p53 when transcription is found to be the same
CC or lower in the first, than in s2. The methods and products can be used
CC to determine p53 status, to diagnose cancer and to evaluate cytotoxicity
CC or carcinogenicity of a test agent. X86265-75 encode human PIG proteins.
XX
SQ Sequence 2302 BP; 497 A; 752 C; 594 G; 459 T; 0 other;

Query Match 48.0%; Score 19.2; DB 20; Length 2302;
Best Local Similarity 75.0%; Pred. No. 37;
Matches 24; Conservative 0; Mismatches 8; Indels 0; Gaps 0;
Qy 1 ctcgagccaccataaccctcaatactccag 32
   ||| ||| ||| ||| ||| ||| ||| |||
Db 42 CTCGACTTAGCCCACTACTACCTCATGACTCCAG 11

RESULT 13
Q50146
ID Q50146 standard; DNA; 2401 BP.
XX
```

```
AC Q50146;
XX
DT 03-JUN-1994 (first entry)
XX
DE Restriction enzyme.
XX
KW Restriction site; enzyme; SphI; BamHI; cleavage-site;
KW phage; Escherichia coli; ss.
XX
Key :Location/Qualifiers
FH mat_peptide 240..1074
FT /*tag= b
FT misc_feature 897
FT /*tag= c
FT /*label= SphI_cleavage-site
FT mat_peptide 1127..2008
FT /*tag= d
FT misc_feature 1425
FT /*tag= e
FT /*label= BamHI_cleavage-site
XX
PN JP05244946-A.
XX
PD 24-SEP-1993.
XX
PF 03-MAR-1992; 92JP-0094990.
XX
PR 03-MAR-1992; 92JP-0094990.
XX
PA (KAJI/) KAJI A..
XX
DR WPI; 1993-389087/49.
DR P-PSDB; R42305.
XX
PT New restriction enzyme - suppresses melting by T-even phage of
PT Escherichia coli
XX
PS Claim 1; Page 7-10; 10pp; Japanese.
XX
CC The enzyme can suppress the melting by T-even phage of E.coli, in a
CC temp. dependent manner. A large amt. of enzyme can be produced.
XX
SQ Sequence 2401 BP; 681 A; 564 C; 543 G; 613 T; 0 other;

Query Match 47.5%; Score 19; DB 14; Length 2401;
Best Local Similarity 71.4%; Pred. No. 46;
Matches 25; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
Qy 3 ctagccaccataaccctcaatactccaggatt 37
   ||| ||| ||| ||| ||| ||| ||| |||
Db 284 caaggcaccgcacacacgcaatacacacaggcgatt 318

RESULT 14
X14116
ID X14116 standard; DNA; 1183 BP.
XX
AC X14116;
XX
DT 31-MAR-1999 (first entry)
XX
DE H. pylori GHPO 106 gene.
XX
KW GHPO protein; Helicobacter infection; gastroduodenal disease; gastritis;
KW peptic ulcer disease; ss.
XX
OS Helicobacter pylori.
XX
FH Key :Location/Qualifiers
FT CDS 51..1133
FT /*tag= a
XX
```

```
PN WO9843478-A1.
XX
PD 08-OCT-1998.
XX
PF 01-APR-1998; 98WO-US06371.
XX
PR 29-JUL-1997; 97US-0902615.
PR 01-APR-1997; 97US-0833457.
PR 24-JUN-1997; 97US-0881227.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA (INMR ) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS.
XX
PI Al-Garawi A, Kleanthous H, Miller C, Oomen RP, Tomb J;
XX WPI; 1998-542293/46.
DR P-PSDB; W98397.
XX
XX New isolated Helicobacter polynucleotides - used to develop products
PT for the diagnosis, prevention and treatment of Helicobacter
PT infections and gastrointestinal diseases
XX
XX Claim 1; Page 650-651; 2054pp; English.
XX
XX This sequence represents a polynucleotide of the invention. It was
CC isolated from Helicobacter pylori and encodes a H.pylori GHP0 protein.
CC The polypeptides can be used for preventing or treating Helicobacter
CC infections, and gastroduodenal diseases associated with these
CC infections, including acute, chronic, and atrophic gastritis, and peptic
CC ulcer diseases, e.g. gastric and duodenal ulcers. They can also be used
CC for the production of antibodies. The products can also be used for
CC detection and diagnosis.
XX
SQ Sequence 1183 BP; 413 A; 200 C; 242 G; 328 T; 0 other;

Query Match 47.0%; Score 18.8; DB 19; Length 1183;
Best Local Similarity 68.4%; Pred. No. 49;
Matches 26; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

QY 2 tcgacccaccataaccctcaatactccaggattgg 39
Db ||||| | ||||| ||||| ||||| |||||
355 tcgtgagcgtcataaccctcaatactccatgaatgg 392

RESULT 15
292219/C
ID 292219 standard; DNA; 2191 BP.
XX
AC 292219;
XX
XX 05-JUN-2000 (first entry)
XX
XX Human thioredoxin reductase nucleotide sequence #1.
XX
XX Thioredoxin reductase; redox inhibitor; cellular proliferation; cancer;
KW apoptosis; reperfusion injury; hepatitis; amyotrophic lateral sclerosis;
KW Alzheimer's disease; autoimmune disease; Sjogren's syndrome; lupus;
KW rheumatoid arthritis; HIV; Hermansky-Pudlak syndrome; retinopathy;
KW wound healing; Crohn's disease; ulcerative colitis; angiogenesis;
KW adult respiratory distress syndrome; ARDS; infection; ds;
KW inflammatory conditions; psoriasis; inflammatory bowel disease;
KW hyperproliferative disease; mycosis fungoides.
XX
OS Homo sapiens.
XX
XX WO200006088-A2.
XX
PD 10-FEB-2000.
XX
PF 02-AUG-1999; 99WO-US17496.
XX
PR 31-JUL-1998; 98US-0127219.
```

```
XX (KIRK/) KIRKPATRICK D L.
PA (POWI/) POWIS G.
XX
XX Kirkpatrick DL; Powis G;
XX
XX WPI; 2000-195158/17.
XX
XX New composition for preventing inhibition of apoptosis and for treating
PT e.g. cancer or reperfusion injuries comprises inhibitor of cellular
PT redox signalling
XX
XX Disclosure; Page 51-53; 55pp; English.
XX
XX This sequence represents a thioredoxin reductase nucleotide sequence.
CC Thioredoxin exerts specific redox control over a number of transcription
CC factors to modulate their DNA binding and therefore regulates gene
CC transcription. Thioredoxin reductase catalyses the reduction of
CC thioredoxin. The thioredoxin redox system plays an important role in a
CC number of diseased states. The invention relates to a composition
CC comprising an inhibitor of cellular redox signalling and a carrier. The
CC composition can be used for inhibiting cellular proliferation or
CC preventing inhibition of apoptosis and is therefore useful for treating
CC cancer, reperfusion injury following ischaemia, hepatitis, amyotrophic
CC lateral sclerosis, neurodegenerative diseases, Alzheimer's disease,
CC autoimmune diseases, Sjogren's syndrome, lupus, rheumatoid arthritis,
CC HIV, Hermansky-Pudlak syndrome, retinal oxidative damage, retinopathy,
CC skin hyperplasia, aging, ultraviolet damage, wound healing, Crohn's
CC disease, ulcerative colitis, angiogenesis, uterine disorders, adult
CC respiratory distress syndrome (ARDS), lung disorders, viral infections
CC such as herpes virus, pox virus and adenovirus infections, inflammatory
CC conditions, autoimmune diseases (such as systemic lupus erythematosus,
CC rheumatoid arthritis, psoriasis, inflammatory bowel disease and
CC autoimmune diabetes), immune mediated glomerular nephritis,
CC hyperproliferative diseases (such as fibrosis) and mycosis fungoides.
XX
SQ Sequence 2191 BP; 614 A; 411 C; 555 G; 611 T; 0 other;

Query Match 47.0%; Score 18.8; DB 21; Length 2191;
Best Local Similarity 76.7%; Pred. No. 55;
Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 5 agccacccataaccctcaatactccagg 34
Db ||||| | ||||| ||||| |||||
878 AGCCAGCCAGAACCCCTGCACACTCCAGG 849

Search completed: January 18, 2001, 11:04:27
Job time: 29436 sec
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pct-us00-32259-12.rng

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C 2	23	57.5	2634	2	US-08-818-514-2		Sequence 2, Appli
C 3	23	57.5	2634	3	US-09-115-934A-1		Sequence 1, Appli
C 4	23	57.5	2634	3	US-09-115-934A-2		Sequence 2, Appli
C 5	20	50.0	3364	2	US-08-735-609-9		Sequence 9, Appli
C 6	20	50.0	3364	2	US-08-735-609-9		Sequence 9, Appli
C 7	20	50.0	3364	3	US-09-315-372-9		Sequence 9, Appli
C 8	20	50.0	3364	3	US-09-245-752-9		Sequence 9, Appli
C 9	20	50.0	3364	3	US-09-245-497-9		Sequence 9, Appli
C 10	19.2	48.0	1864	3	US-09-221-235-10		Sequence 10, Appli
C 11	19.2	48.0	1864	3	US-09-221-928-10		Sequence 10, Appli
C 12	19.2	48.0	1864	3	US-09-221-527-10		Sequence 10, Appli
C 13	19.2	48.0	1864	3	US-09-221-236-10		Sequence 10, Appli
C 14	19.2	48.0	1864	3	US-09-221-416-10		Sequence 10, Appli
C 15	18.6	47.0	3207	2	US-08-770-379-20		Sequence 20, Appli
C 16	18.6	46.5	798	2	US-08-743-009-2		Sequence 2, Appli
C 17	18.6	46.5	798	3	US-09-172-110-2		Sequence 2, Appli
C 18	18.4	46.0	1045	1	US-08-317-880-3		Sequence 3, Appli
C 19	18.4	46.0	1045	2	US-08-783-396-3		Sequence 3, Appli
C 20	18.4	46.0	2425	2	US-08-132-990A-3		Sequence 3, Appli
C 21	18.4	46.0	2425	4	PCR-US92-09382-3		Sequence 3, Appli
C 22	18.4	46.0	3954	1	US-08-317-880-1		Sequence 1, Appli
C 23	18.4	46.0	3954	2	US-08-783-396-1		Sequence 1, Appli
C 24	18.2	45.5	259	2	US-08-493-137-3		Sequence 3, Appli
C 25	18.2	45.5	259	4	PCR-US95-07844-3		Sequence 3, Appli
C 26	18.2	45.5	1305	3	US-09-013-072-3		Sequence 3, Appli
C 27	18.2	45.5	2584	3	US-08-758-662-8		Sequence 8, Appli
C 28	18.2	45.5	2784	1	US-08-474-140-10		Sequence 10, Appli



Hartigan-O'Connor, Dennis J.  
TITLE OF INVENTION: IMPROVED ADENOVIRUS VECTORS  
NUMBER OF SEQUENCES: 15  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Medlen & Carroll, LLP  
STREET: 220 Montgomery Street, Suite 2200  
CITY: San Francisco  
STATE: California  
COUNTRY: United States Of America  
ZIP: 94104

ZIP: 94104  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: Patentin Release #1.0, Version #1.30  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/735,609  
 FILING DATE: 23-Oct-1996  
 CLASSIFICATION: <unknown>

CLASSIFICATION: <UNKNOWN>  
ATTORNEY/AGENT INFORMATION:  
NAME: Ingolia, Diane E.  
REGISTRATION NUMBER: 40-027  
REFERENCE/DOCKET NUMBER: UM-02484  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 705-8410  
TELEFAX: (415) 397-8338  
FORMATION FOR SEQ ID NO: 9:

SEQUENCE CHARACTERISTICS:  
LENGTH: 3364 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "DNA"  
SEQUENCE DESCRIPTION: SEQ ID NO: 9:  
1-609-9

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Match          50.08; Score 20; DB 2; Length 3364;
Local Similarity 72.28; Pred. No. 11;
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      |||||
      |||||
      |||||
      AGCCTGCCCCCTCACCTGGATTCACAGGACAGGG 2291

```

MICANT: Hauser, Michael A.  
 MICANT: Kumar-Singh, Rajendra  
 MICANT: Hartigan-O'Connor, Dennis J.  
 DATE OF INVENTION: IMPROVED ADENOVIRUS VECTORS  
 NUMBER OF SEQUENCES: 15  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Medlen & Carroll, LLP  
 STREET: 220 Montgomery Street, Suite 2200  
 CITY: San Francisco  
 STATE: California  
 COUNTRY: United States Of America  
 ZIP: 94104

COUNTRY: United States Of America  
CITY: California  
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WWW: WWW.PACIFIC.COM  
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SUITE 100  
SAN FRANCISCO CA 94104  
UNITED STATES OF AMERICA

FLOPPY DISK  
IBM PC COMPATIBLE  
PC-DOS/MG-DOS  
PATENT IN RELEASE #1.0, VERSION #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/315.372

; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/735,609  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Ingolia, Diane E.  
; REGISTRATION NUMBER: 40,027  
; REFERENCE/DOCKET NUMBER: UM-02484  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 705-8410  
; TELEFAX: (415) 397-8338  
; INFORMATION FOR SEQ ID NO: 9:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 3364 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: other nucleic acid  
; DESCRIPTION: /desc = "DNA"  
US-09-315-372-9

Query Match 50.0%; Score 20; DB 3; Length 3364;  
Best Local Similarity 72.2%; Pred. No. 11;  
Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 5 agccaccctaacccctcaatactccaggattggg 40  
||||| ||||| ||||| ||||| ||||| ||||| |||||  
Db 2326 AGCGTCCCTCACCCCTGGATCCACGAGGACAGG 2291

RESULT 8  
US-09-244-752-9/c  
; Sequence 9, Application US/09244752  
; Patent No. 6063622  
; GENERAL INFORMATION:  
; APPLICANT: Chamberlain, Jeffrey S.  
; APPLICANT: Amalfitano, Andrea  
; APPLICANT: Hauser, Michael A.  
; APPLICANT: Kumar-Singh, Rajendra  
; APPLICANT: Hartigan-O'Connor, Dennis J.  
; TITLE OF INVENTION: IMPROVED ADENOVIRUS VECTORS  
; NUMBER OF SEQUENCES: 15  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Medlen & Carroll, LLP  
; STREET: 220 Montgomery Street, Suite 2200  
; CITY: San Francisco  
; STATE: California  
; COUNTRY: United States Of America  
; ZIP: 94104  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
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; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/244,752  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/735,609  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Ingolia, Diane E.  
; REGISTRATION NUMBER: 40,027  
; REFERENCE/DOCKET NUMBER: UM-02484  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 705-8410  
; TELEFAX: (415) 397-8338  
; INFORMATION FOR SEQ ID NO: 9:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 3364 base pairs

; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: other nucleic acid  
; DESCRIPTION: /desc = "DNA"  
US-09-244-752-9

Query Match 50.0%; Score 20; DB 3; Length 3364;  
Best Local Similarity 72.2%; Pred. No. 11;  
Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 5 agccaccctaacccctcaatactccaggattggg 40  
||||| ||||| ||||| ||||| ||||| ||||| |||||  
Db 2326 AGCGTCCCTCACCCCTGGATCCACGAGGACAGG 2291

RESULT 9  
US-09-245-497-9/c  
; Sequence 9, Application US/09245497  
; Patent No. 6083750  
; GENERAL INFORMATION:  
; APPLICANT: Chamberlain, Jeffrey S.  
; APPLICANT: Amalfitano, Andrea  
; APPLICANT: Hauser, Michael A.  
; APPLICANT: Kumar-Singh, Rajendra  
; APPLICANT: Hartigan-O'Connor, Dennis J.  
; TITLE OF INVENTION: IMPROVED ADENOVIRUS VECTORS  
; NUMBER OF SEQUENCES: 15  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Medlen & Carroll, LLP  
; STREET: 220 Montgomery Street, Suite 2200  
; CITY: San Francisco  
; STATE: California  
; COUNTRY: United States Of America  
; ZIP: 94104  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/245,497  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/735,609  
; APPLICATION NUMBER: <B> FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Ingolia, Diane E.  
; REGISTRATION NUMBER: 40,027  
; REFERENCE/DOCKET NUMBER: UM-02484  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 705-8410  
; TELEFAX: (415) 397-8338  
; INFORMATION FOR SEQ ID NO: 9:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 3364 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: other nucleic acid  
; DESCRIPTION: /desc = "DNA"  
US-09-245-497-9

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Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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Db 2326 AGCGTCCCTCACCCCTGGATCCACGAGGACAGG 2291

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; APPLICANT: Acton,, Susan
; TITLE OF INVENTION: NOVEL CSAPK-1 NUCLEIC ACID MOLECULES AND USES THEREFOR
; FILE REFERENCE: MNI-050
; CURRENT APPLICATION NUMBER: US/09/221,527
; CURRENT FILING DATE: 1998-12-28
; EARLIER APPLICATION NUMBER: 09/163,115
; EARLIER FILING DATE:
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 10
; LENGTH: 1864
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (275)..(754)
US-09-221-527-10
;
Query Match      48.0%; Score 19.2; DB 3; Length 1864;
Best Local Similarity 67.5%; Pred. No. 21;
Matches 27; Conservative 0; Mismatches 13; Indels 0; Gaps 0

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      ||| ||| |||| | ||| |||| | |||| | ||||
Db    955 ctccagcattccccaaagtcttaattcccataaatggg 994

RESULT 13
US-09-221-236-10
; Sequence 10, Application US/09221236
; Patent No. 6146841
; GENERAL INFORMATION:
; APPLICANT: Acton,, Susan
; TITLE OF INVENTION: NOVEL CSAPK-1 NUCLEIC ACID MOLECULES AND USES THEREFOR
; FILE REFERENCE: MNI-050
; CURRENT APPLICATION NUMBER: US/09/221,236
; CURRENT FILING DATE: 1998-12-28
; EARLIER APPLICATION NUMBER: 09/163,115
; EARLIER FILING DATE: 1998-09-29
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 10
; LENGTH: 1864
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (275)..(754)
US-09-221-236-10
;
Query Match      48.0%; Score 19.2; DB 3; Length 1864;
Best Local Similarity 67.5%; Pred. No. 21;
Matches 27; Conservative 0; Mismatches 13; Indels 0; Gaps 0

QY   1 ctcgagccaccataacccctcaataactcaggattggg 40
      ||| ||| |||| | ||| |||| | |||| | ||||
Db    955 ctccagcattccccaaagtcttaattcccataaatggg 994

RESULT 14
US-09-221-416-10
; Sequence 10, Application US/09221416
; Patent No. 6153417
; GENERAL INFORMATION:
; APPLICANT: Acton,, Susan
; TITLE OF INVENTION: NOVEL CSAPK-1 NUCLEIC ACID MOLECULES AND USES THEREFOR
; FILE REFERENCE: MNI-050
; CURRENT APPLICATION NUMBER: US/09/221,416
; CURRENT FILING DATE: 1998-12-28
; EARLIER APPLICATION NUMBER: 09/163,115
; EARLIER FILING DATE: 1998-09-29

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; NUMBER OF SEQ ID NOS: 15  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 10  
; LENGTH: 1864  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (275)..(754)  
US-09-221-416-10

Query Match 48.0%; Score 19.2; DB 3; Length 1864;  
Best Local Similarity 67.5%; Pred. No. 21;  
Matches 27; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

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RESULT 15

US-08-770-379-20  
; Sequence 20, Application US/08770379  
; Patent No. 5849564  
; GENERAL INFORMATION:  
; APPLICANT: Chang, Yuan  
; APPLICANT: Bohenzky, Roy A.  
; APPLICANT: Russo, James J.  
; APPLICANT: Edelman, Isidore S.  
; APPLICANT: Moore, Patrick S.  
; TITLE OF INVENTION: POLYPEPTIDES FROM KAPOSI'S SARCOMA-ASSOCIATED  
; TITLE OF INVENTION: HERPESVIRUS, DNA ENCODING SAME AND USES THEREOF  
; NUMBER OF SEQUENCES: 20  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Cooper & Dunham LLP  
; STREET: 1185 Avenue of the Americas  
; CITY: New York  
; STATE: New York  
; COUNTRY: U.S.A.  
; ZIP: 10036  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/770,379  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: White, John P.  
; REGISTRATION NUMBER: 28,678  
; REFERENCE/DOCKET NUMBER: 52342  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (212) 278-0400  
; TELEFAX: (212) 391-0525  
; INFORMATION FOR SEQ ID NO: 20:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 32207 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-770-379-20

Query Match 47.0%; Score 18.8; DB 2; Length 32207;  
Best Local Similarity 76.7%; Pred. No. 54;  
Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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Db 3464 ACCATATCACCATCCACACTCCAGGGACTG 3493

Search completed: January 18, 2001, 10:59:56  
Job time: 30474 sec





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GenCore version 4.5  
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model  
Run on: January 18, 2001, 03:53:45 ; Search time 3453.92 Seconds  
(without alignments)  
81.154 Million cell updates/sec

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Perfect score: 40  
Sequence: 1 ctcgagccaccctcaataccctcaatccaggattggg 40

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 7991742 seqs, 3503743858 residues  
Total number of hits satisfying chosen parameters: 15983484

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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193: gb\_gss28:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	24.6	61.5	296	96	AW845395	AW845395 CM4-CT004
C 2	23.8	59.5	250	145	T08487	T08487 EST06378 In
C 3	23.6	59.0	337	146	T92497	T92497 Y024909.r1
C 4	23.5	57.5	69	137	BE878784	BE878784 601493062
C 5	23.5	57.5	118	95	AW804230	AW804230 PM3-UM008
C 6	23.5	57.5	135	133	BE005490	BE005490 CM1-BN011
C 7	23.5	57.5	147	133	BE005492	BE005492 CM1-BN011
C 8	23.5	57.5	161	143	N73434	N73434 Y231e12.r1
C 9	23.5	57.5	223	134	BE089733	BE089733 QV0-BT070
C 10	23.5	57.5	227	133	BE005486	BE005486 CM1-BN011
C 11	23.5	57.5	231	5	AA327421	AA327421 EST30756
C 12	23.5	57.5	237	139	BF037734	BF037734 601462081
C 13	23.5	57.5	249	95	AW804233	AW804233 PM3-UM008
C 14	23.5	57.5	251	110	BE622136	BE622136 601440862
C 15	23.5	57.5	252	136	BE869246	BE869246 601445061
C 16	23.5	57.5	252	137	BE894967	BE894967 601436172
C 17	23.5	57.5	260	2	AA101534	AA101534 zn79c11.s
C 18	23.5	57.5	263	6	AA383010	AA383010 EST96488
C 19	23.5	57.5	265	137	BE878754	BE878754 601493017
C 20	23.5	57.5	276	110	BE617770	BE617770 601441968
C 21	23.5	57.5	289	145	T39447	T39447 Y066508.r1
C 22	23.5	57.5	299	104	BE178852	BE178852 PM1-HT060
C 23	23.5	57.5	304	6	AA384401	AA384401 EST97914
C 24	23.5	57.5	305	12	AA853875	AA853875 NHT8Cae09
C 25	23.5	57.5	307	95	AW804241	AW804241 PM3-UM008
C 26	23.5	57.5	307	145	T39458	T39458 Y066507.r2
C 27	23.5	57.5	308	6	AA383479	AA383479 EST96859
C 28	23.5	57.5	308	95	AW804238	AW804238 PM3-UM008
C 29	23.5	57.5	313	137	BE936437	BE936437 RC1-NT003
C 30	23.5	57.5	326	6	AA361993	AA361993 EST71440
C 31	23.5	57.5	328	95	AW804271	AW804271 PM3-UM008
C 32	23.5	57.5	329	135	BE763332	BE763332 RC1-NT003
C 33	23.5	57.5	332	6	AA373764	AA373764 EST85818
C 34	23.5	57.5	334	144	R66220	R66220 Y134807.r1
C 35	23.5	57.5	335	145	T52898	T52898 Y081112.r1
C 36	23.5	57.5	337	38	AV703130	AV703130 AV703130
C 37	23.5	57.5	342	6	AA363971	AA363971 EST74461
C 38	23.5	57.5	343	137	BE936435	BE936435 RC1-NT003
C 39	23.5	57.5	344	95	AW804222	AW804222 PM3-UM008
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C 41	23.5	57.5	348	134	BE089688	BE089688 QV0-BT070
C 42	23.5	57.5	349	134	BE089689	BE089689 QV0-BT070
C 43	23.5	57.5	353	133	BE003833	BE003833 QV3-BN009
C 44	23.5	57.5	354	96	AW904630	AW904630 RC1-NN106
C 45	23.5	57.5	361	133	BE005530	BE005530 CM1-BN011

## ALIGNMENTS

RESULT 1  
AW845395/c  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

AW845395 296 bp mRNA EST  
CM4-CT0045-180200-512-a09 CT0045 Homo sapiens cdna, mRNA sequence.  
AW845395  
AW845395.1 GI:7940912  
EST.  
human.  
Homo sapiens







source

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/organism="Homo sapiens"
/db_xref="GDB:3903214"
/db_xref="taxon:9606"
/clone="IMAGE:284686"
/clone_lib="Soares_multiple_sclerosis_2NBHMS"
/sex="male"
/tissue_type="multiple sclerosis lesions"
/dev_stage="Age 46"
/lab_host="DH10B (ampicillin resistant)"
/notes="Vector: pMT73D (Pharmacia) with a modified
polylinker V-type: phagemid; Site_1: Not I; Site_2: Eco RI
; 1st strand cDNA was primed with a Not I - oligo(dT)
primer [5',
TGTACCAATCTGAAGTGGGAGCGCGCCATTTTTTTTTTTTTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pMT73 vector
(Pharmacia). Library went through one round of
normalization to a Cot = 5. Library constructed by Bento
Soares and M.Patima Bonaldo. RNA from 4 multiple sclerosis
lesions from one patient was kindly provided by Dr. Kevin
G. Becker (NINDS/NIH)."
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BASE COUNT 36 a 45 c 44 g 36 t

ORIGIN

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Query Match 57.5%; Score 23; DB 143; Length 161;
Best Local Similarity 74.4%; Pred. No. 37;
Matches 29; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
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Qy 2 tcgagccaccataaccctcaatactccaggattggg 40

Db 91 TCACCCACACTTGAAGCCACAAACTGCAGGATTGGG 53

RESULT 9

BE005486

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

COMMENT

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome

Project. This entry can be seen in the following URL

(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=et2-QV0-BT0704-120

500-224-c08&t3-2000-05-12&t4=1)

Seq primer: puc 18 forward

High quality sequence start: 12

High quality sequence stop: 223.

FEATURES

source

1. .223

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/db\_xref="taxon:9606"

/clone\_lib="BT0704"

/dev\_stage="Adult"

/note="Organ: breast; Vector: puc18; Site\_1: SmaI; Site\_2:

SmaI; A mini-library was made by cloning products derived

from ORESTES PCR (U.S. Letters Patent application No. 196

,716 - Ludwig Institute for Cancer Research) profiles

into the puc 18 vector. Reverse transcription of tissue

mRNA and cDNA amplification were performed under low

stringency conditions."

BASE COUNT 47 a 52 c 70 g 54 t

ORIGIN

Query Match 57.5%; Score 23; DB 134; Length 223;

Best Local Similarity 74.4%; Pred. No. 38;

Matches 29; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 2 tcgagccaccataaccctcaatactccaggattggg 40

Db 86 TCACCCACACTTGAAGCCACAAACTGCAGGATTGGG 48

RESULT 10

BE005486

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

COMMENT

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome

Project. This entry can be seen in the following URL

(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=et2-CM1-BN0117-110

400-183-e09&t3-2000-04-11&t4=1)

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High quality sequence stop: 227.

FEATURES

source

1. .227

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/db\_xref="taxon:9606"

/clone\_lib="BN0117"

/dev\_stage="Adult"

/note="Organ: breast\_normal; Vector: puc18; Site\_1: SmaI;

Site\_2: SmaI; A mini-library was made by cloning products

derived from ORESTES PCR (U.S. Letters Patent application

No. 196,716 - Ludwig Institute for Cancer Research)









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